

submitted its comment and petition. The comment, therefore, appears to be unrelated and inapplicable to the instant action.

The petition also seems to allege that, because the FAA employs standardized language in its routine airspace action rulemakings, it has not performed the requisite analyses, specifically with respect to the Regulatory Flexibility Act and Executive Order (E.O.) 12866. Indeed, the FAA has developed standardized language for use in these routine actions to ensure consistency and streamline drafting of rulemaking documents in one of its most prolific regulatory portfolios. However, each action is individually assessed and tailored as appropriate in compliance with applicable law and agency guidance, including the Administrative Procedures Act; the Regulatory Flexibility Act; E.O. 12866; FAA Order JO 7400.2R, *Procedures for Handling Airspace Matters*; and FAA Order 1050.1G, *FAA National Environmental Policy Act Implementing Procedures*. Ironically, the commenter appears to utilize boilerplate language in its comment and petition that is not relevant to the instant action.

Substantively, the petition also makes an assumption that the expansion of the Class E airspace extending upward from 700 ft. above the surface at Beeville, TX, “can foreclose or constrain marginal VFR operations,” which is unsubstantiated. This Class E airspace is established as transitional airspace to/ from terminal or enroute environments around airports with instrument procedures as required by FAA Order JO 7400.2R. No air traffic control services are provided, and there are no communications requirements, so there is little to no impact on VFR traffic. In fact, this Class E airspace advises the VFR pilots that there are instrument procedures and potential IFR traffic in the area which increases the pilot’s situational awareness.

Incorporation by Reference

Class E airspace designations are published in paragraph 6005 of FAA Order JO 7400.11, *Airspace Designations and Reporting Points*, which is incorporated by reference in 14 CFR 71.1 on an annual basis. This document amends the current version of that order, FAA Order JO 7400.11K, dated August 4, 2025, and effective September 15, 2025. These amendments will be published in the next update to FAA Order JO 7400.11. FAA Order JO 7400.11K, which lists Class A, B, C, D, and E airspace areas, air traffic service routes, and reporting points, is publicly

available as listed in the **ADDRESSES** section of this document.

The Rule

This action amends 14 CFR part 71 by modifying the Class E airspace extending upward from 700 ft. above the surface at Beeville, Texas, due to an airspace review conducted as part of the decommissioning of the Three Rivers VOR as part of the VOR MON Program.

For the Beeville Municipal Airport, Beeville, TX, Class E airspace extending upward from 700 ft. above the surface, this action: (1) increases the radius from 6.6 miles to 7.1 miles from the airport; (2) removes the Beeville NDB and associated extension as they are no longer required; and (3) updates the geographic coordinates of the airport to coincide with the FAA’s aeronautical database.

Regulatory Notices and Analyses

The FAA has determined that this regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. It, therefore: (1) is not a “significant regulatory action” under Executive Order 12866; (2) is not a “significant rule” under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. Since this is a routine matter that only affects air traffic procedures and air navigation, it is certified that this rule, when promulgated, does not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

Environmental Review

The FAA has determined that this action qualifies for categorical exclusion under the National Environmental Policy Act in accordance with FAA Order 1050.1G, “FAA National Environmental Policy Act Implementing Procedures,” Paragraph B–2.5(a). This airspace action is not expected to cause any potentially significant environmental impacts, and no extraordinary circumstances exist that warrant preparation of an environmental assessment.

Lists of Subjects in 14 CFR 71

Airspace, Incorporation by reference, Navigation (air).

The Amendment

In consideration of the foregoing, the Federal Aviation Administration amends 14 CFR part 71 as follows:

PART 71—DESIGNATION OF CLASS A, B, C, D, AND E AIRSPACE AREAS; AIR TRAFFIC SERVICE ROUTES; AND REPORTING POINTS

■ 1. The authority citation for 14 CFR part 71 continues to read as follows:

Authority: 49 U.S.C. 106(f), 106(g), 40103, 40113, 40120; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

§ 71.1 [Amended]

■ 2. The incorporation by reference in 14 CFR 71.1 of FAA Order JO 7400.11K, *Airspace Designations and Reporting Points*, dated August 4, 2025, and effective September 15, 2025, is amended as follows:

Paragraph 6005 Class E Airspace Areas Extending Upward From 700 Feet or More Above the Surface of the Earth.

* * * * *

ASW TX E5 Beeville, TX [Amended]

Beeville Municipal Airport
(Lat. 28°21’51” N, long. 097°47’31” W)

That airspace extending upward from 700 feet above the surface within a 7.1-mile radius of Beeville Municipal Airport.

* * * * *

Issued in Fort Worth, Texas, on February 4, 2026.

Courtney E. Johns,

*Acting Manager, Operations Support Group,
ATO Central Service Center.*

[FR Doc. 2026–02319 Filed 2–5–26; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF ENERGY

Federal Energy Regulatory Commission.

18 CFR Part 157

[Docket No. RM25–14–000]

Implementation of the Executive Order Entitled “Zero-Based Regulatory Budgeting To Unleash American Energy”; Correction

AGENCY: Federal Energy Regulatory Commission.

ACTION: Correcting amendment.

SUMMARY: The Federal Energy Regulatory Commission (FERC) published a direct final rule in the **Federal Register** of October 21, 2025, revising its regulations to insert a conditional sunset date into certain regulations in response to Executive Order 14270, “Zero-Based Regulatory Budgeting to Unleash American Energy.” The document contained an error. This document corrects the regulations.

DATES: This correction is effective February 6, 2026.

FOR FURTHER INFORMATION CONTACT: Karin Herzfeld, Office of the General Counsel, Federal Energy Regulatory Commission, 888 First Street NE, Washington, DC 20426, (202) 502-8459, karin.herzfeld@ferc.gov.

SUPPLEMENTARY INFORMATION: In FR Doc. 2025-19607 (193 FERC ¶ 61,002) (90 FR 48397, October 21, 2025), FERC added a conditional sunset date to § 157.202 at paragraph (b)(2)(ii)(H) in error. FERC is removing paragraph (b)(2)(ii)(H).

List of Subject in 18 CFR Part 157

Administrative practice and procedure, Natural gas, Reporting and recordkeeping requirements.

Accordingly, 18 CFR part 157 is corrected by making the following correcting amendment:

PART 157—APPLICATIONS FOR CERTIFICATES OF PUBLIC CONVENIENCE AND NECESSITY AND FOR ORDERS PERMITTING AND APPROVING ABANDONMENT UNDER SECTION 7 OF THE NATURAL GAS ACT

■ 1. The authority citation for part 157 continues to read as follows:

Authority: 15 U.S.C. 717-717w, 3301-3432; 42 U.S.C. 7101-7352.

§ 157.202 [Amended]

■ 2. In § 157.202, remove paragraph (b)(2)(ii)(H).

Issued: February 4, 2026.

Debbie-Anne A. Reese,
Secretary.

[FR Doc. 2026-02431 Filed 2-5-26; 8:45 am]

BILLING CODE 6717-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 73

[Docket No. FDA-2024-C-3384]

Listing of Color Additives Exempt From Certification; Spirulina Extract

AGENCY: Food and Drug Administration, HHS.

ACTION: Final amendment; order.

SUMMARY: The Food and Drug Administration (FDA or we) is amending the color additive regulations to provide for the expanded use of spirulina (*Arthrospira platensis*) extract as a color additive in human foods generally (except for infant formula,

certain foods subject to regulation by the U.S. Department of Agriculture, and foods for which standards of identity have been issued under section 401 of the Federal Food, Drug, and Cosmetic Act, unless the use of the added color is authorized by such standards) at levels consistent with good manufacturing practice (GMP), to lower the heavy metal specifications for lead, arsenic, and mercury, and to add a specification for cadmium. We are taking this action in response to a color additive petition (CAP) submitted by GNT USA, LLC (GNT or petitioner).

DATES: This order is effective March 23, 2026. See section IX of this document for further information on the filing of objections. Either electronic or written objections and requests for a hearing on the order must be submitted by March 9, 2026.

ADDRESSES: You may submit objections and requests for a hearing as follows. Please note that late, untimely filed objections will not be considered. The <https://www.regulations.gov> electronic filing system will accept objections until 11:59 p.m. Eastern Time at the end of March 9, 2026. Objections received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic objections in the following way:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the instructions for submitting comments. Objections submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your objection will be made public, you are solely responsible for ensuring that your objection does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your objection, that information will be posted on <https://www.regulations.gov>.

- If you want to submit an objection with confidential information that you do not wish to be made available to the public, submit the objection as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- **Mail/Hand Delivery/Courier (for written/paper submissions):** Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper objections submitted to the Dockets Management Staff, FDA will post your objection, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA-2024-C-3384 for "Listing of Color Additives Exempt from Certification; Spirulina Extract." Received objections, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <https://www.regulations.gov> or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240-402-7500.

- **Confidential Submissions—**To submit an objection with confidential information that you do not wish to be made publicly available, submit your objections only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." We will review this copy, including the claimed confidential information, in our consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <https://www.regulations.gov>.